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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
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10/080,043

02/22/2002

Oliver Yoa-Pu Hu

26193-167918

8602

38598

7590

10/11/2006

ANDREWS KURTH LLP
1350 I STREET, N.W.
SUITE 1100
WASHINGTON, DC 20005

EXAMINER

SPIVACK, PHYLLIS G

ART UNIT

PAPER NUMBER

1614

DATE MAILED: 10/11/2006

Please find below and/or attached an Office communication concerning this application or proceeding.



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There was no overpayment made by applicant. All fees were calculated and assessed properly. The fees were charged to the deposit account on the document at the time of submission although the fee was not deducted from the account until November 2005.

Sincerely,

A handwritten signature in cursive script, appearing to read "Deborah E. Dotson", is written over the typed name.

Deborah E. Dotson
Technical Center 1600
571 272 0520



FILE COPY

UNITED STATES PATENT AND TRADEMARK OFFICE

UNDER SECRETARY OF COMMERCE FOR INTELLECTUAL PROPERTY AND
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Sincerely,

Deborah E. Dotson
Technical Center 1600
571 272 0520

- call Attorney

ATTENTION ATTENTION ATTENTION

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Patent/TM/App/Serial # 10,080,043

Program Area Leek Center 1614

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ATTENTION ATTENTION ATTENTION



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

PATENT MAINTENANCE
DIVISION

In re application of:

Hu et al.

Appln. No. 10/080,043

Confirmation No. 8602

Filed: 22 February 2002

For: CYTOCHROME P450 3A INHIBITORS AND
ENHANCERS

Unit: 1614

Examiner: Phyllis G. Spivack

Atty. Docket No. 39297-174169

Customer No.

26694

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REQUEST FOR REFUND

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This is a request for a refund of filing fees, in the amount of \$111.00 (extra claims fees), that the PTO charged to our Deposit Account No. 22-0261 on November 7, 2005. This charge was in error because this file was transferred to another firm in September 2004, and the attorney of record was not authorized to charge on the referenced deposit account at the time the fees were paid. Attached is a copy of the PTO's November 2005 Deposit Account Statement to our firm, highlighting the two charges in question. We do not have any files for this matter and so cannot verify any of the information.

It is respectfully requested that the filing fee, totaling \$111.00, which was charged to this firm's Deposit Account No. 22-0261 on November 7, 2005, be refunded by crediting that amount back to our Deposit Account.

Respectfully submitted,

Michael A. Gollin

Registration No. 31,957

VENABLE LLP

P.O. Box 34385

Washington, D.C. 20043-9998

Phone: (202) 344-4000

FAX: (202) 344-8300

Date: 9/25/06

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11/04	293	11265244	58053-223306	2111	\$250.00	\$61,425.02
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11/07	3	6798526		1562	\$2,300.00	\$56,550.02
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11/07	60	29235826	32405-220144	2112	\$50.00	\$55,764.02
11/07	61	29235826	32405-220144	2312	\$65.00	\$55,699.02
11/07	62	29235826	32405-220144	2051	\$65.00	\$55,634.02

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Email: microberts@venable.com

**FEE TRANSMITTAL
for FY 2003**

Patent fees are subject to annual revision.

☒ Applicant claims small entity status. See 37 CFR 1.27**TOTAL AMOUNT OF PAYMENT (\$)** 55**Complete if Known**

Application Number	10/080,043
Filing Date	February 22, 2002
First Named Inventor	Oliver Yoa-Pu HU et al.
Examiner Name	James H. Reamer
Group / Art Unit	1614
Attorney Docket No.	39297-174169

METHOD OF PAYMENT (check all that apply)☒ Check ☐ Credit card ☐ Money ☐ Other ☐ None
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☐ Charge fee(s) indicated below ☒ Credit any overpayments
☒ Charge any additional fee(s) under 1.18 or 1.17 during pendency of this application
☐ Charge fee(s) indicated below, except for the filing fee to the above-identified deposit account.**FEE CALCULATION****1. BASIC FILING FEE**

Large Entity		Small Entity		Fee Description	Fee Paid
Fee Code	Fee (\$)	Fee Code	Fee (\$)		
1001	750	2001	375	Utility filing fee	
1002	330	2002	165	Design filing fee	
1003	520	2003	260	Plant filing fee	
1004	750	2004	375	Reissue filing fee	
1005	160	2005	80	Provisional filing fee	
SUBTOTAL (1)					(\$) 0

2. EXTRA CLAIM FEES

Total Claims	19	-20 **	=	0	X	Fee from below	=	0
Independent Claims	1	-3 **	=	0	X		=	0
Multiple Dependent					X		=	0

Large Entity		Small Entity		Fee Description	Fee Paid
Fee Code	Fee (\$)	Fee Code	Fee (\$)		
1202	18	2202	9	Claims in excess of 20	
1201	84	2201	42	Independent claims in excess of 3	
1203	280	2203	140	Multiple dependent claim, if not paid	
1204	84	2204	42	** Reissue independent claims over original patent	
1205	18	2205	9	** Reissue claims in excess of 20 and over original patent	
SUBTOTAL (2)					(\$) 0

**or number previously paid, if greater; For Reissues, see above

FEE CALCULATION (continued)**3. ADDITIONAL FEES**

Large Entity		Small Entity		Fee Description	Fee Paid
Fee Code	Fee (\$)	Fee Code	Fee (\$)		
1051	130	2051	65	Surcharge - late filing fee or oath	
1052	50	2052	25	Surcharge - late provisional filing fee or cover sheet	
1053	130	1053	130	Non-English specification	
1812	2,520	1812	2,520	For filing request for ex parte reexam	
1804	820*	1804	820*	Requesting publication of SIR prior to Examiner action	
1805	1,840*	1805	1,840*	Requesting publication of SIR after Examiner action	
1251	110	2251	55	Extension for reply within first month	55
1252	410	2252	205	Extension for reply within second month	
1253	830	2253	465	Extension for reply within third month	
1254	1,450	2254	725	Extension for reply within fourth month	
1255	1,970	2255	985	Extension for reply within fifth month	
1401	320	2401	160	Notice of Appeal	
1402	320	2402	160	Filing a brief in support of an appeal	
1403	280	2403	140	Request for oral hearing	
1451	1,510	1451	1,510	Petition to institute a public use proceeding	
1452	110	2452	55	Petition to revive - unavoidable	
1453	1,300	2453	650	Petition to revive - unintentional	
1501	1,300	2501	650	Utility issue fee (or reissue)	
1502	470	2502	235	Design issue fee	
1503	630	2503	315	Plant issue fee	
1460	130	1460	130	Petitions to the Commissioner	
1807	50	1807	50	Processing fee under 37 CFR 1.17 (a)	
1806	180	1806	180	Submission of Information Disclosure Stmt	
8021	40	8021	40	Recording each patent assignment per property (times number of properties)	
1809	750	2809	375	Filing a submission after final rejection (37 CFR § 1.129(a))	
1810	750	2810	375	For each additional invention to be examined (37 CFR § 1.129(b))	
1801	750	2801	375	Request for Continued Examination (RCE)	
1802	900	1802	900	Request for expedited examination of a design application	
Other fee (specify) _____					
*Reduced by Basic Filing Fee Paid					
SUBTOTAL (3)					(\$) 55

SUBMITTED BY**Complete (if applicable)**

Name (Print/Type)	Fai-Fai Chao, Ph.D.	Registration No. Attorney/Agent	43,538	Telephone	202-344-8011
Signature	Fai-Fai Chao	Date	September 29, 2003		

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Fee History Query

Revenue Accounting and Management

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Total Records Found: 8

Start Date: Any Date

End Date: Any Date

Accounting Date	Sequence Num.	Fee Type	Fee Code	Fee Amount	Mailroom Date	Payment Method
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04/26/2006	00000198	<u>1</u>	<u>2814</u>	\$65.00	04/25/2006	DA 502518
11/07/2005	00000005	<u>1</u>	<u>2202</u>	\$27.00	09/29/2003	DA 220261
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10/27/2005	00000054	<u>1</u>	<u>2253</u>	\$510.00	10/25/2005	DA 502518
10/02/2003	00000121	<u>1</u>	<u>2251</u>	\$55.00	09/29/2003	CK
02/28/2002	00000039	<u>1</u>	<u>201</u>	\$370.00	02/22/2002	CK
02/28/2002	00000040	<u>1</u>	<u>581</u>	\$40.00	02/22/2002	CK

3
2

Sale List

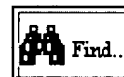
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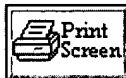
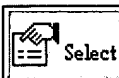
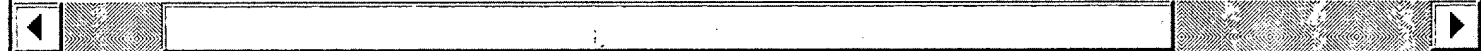


☒ Name/Number:

☐ Attny Docket No:



ing	Operator ID	Seq No.	Sts	Name/Number	Attorney Docket	Dep Act Charge	Other Paymen
6	NNGUYEN1	110	A	10080043	39297-174169	\$125.00	
5	FPATTERS	5	A	10080043	39297-174169	\$111.00	
5	MDAMTE1	22	A	10080043	39297-174169	\$510.00	
8	EFLORES	91	A	10080043	39297-174169		\$55.00
2	SMINASS1	19	A	10080043	39297-174169		\$410.00



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This listing of claims will replace all prior versions, and listings, of claims in the application:

23/5

LISTING OF CLAIMS

41
Claim 1. (Currently Amended) A cytochrome P450 3A (CYP3A) inhibitor ~~which~~
wherein said CYP3A inhibitor is a free base or pharmacologically acceptable salt of at least one
compound selected from the group consisting of α -naphthoflavone, β -naphthoflavone, apigenin,
baicalein, β -myrene, catechin, 3-phenylpropyl acetate, formononetin, ~~gallic acid~~, hesperetin,
~~hesperidin~~, ~~isoquercitrin~~, lauryl alcohol, luteolin, luteolin-7-glycoside, ~~narigin~~,
nordihydroguaiaretic acid, ~~quercitrin~~, and swertiamarin ~~terpineol~~, ~~and trans-cinnamaldehyde~~.

Claim 2. (Cancelled)

Claim 3. (Currently Amended) The CYP3A inhibitor according to claim 1, wherein said
CYP3A inhibitor is at least one selected from the group consisting of nordihydroguaiaretic acid,
(+)-catechin, and lauryl alcohol, ~~gallic acid~~, ~~hesperitin~~, ~~hesperidin~~, ~~trans-cinnamaldehyde~~, β -
~~myrene~~ ~~and narigin~~.

Claim 4. (Canceled)

Claim 5. (Previously Presented) The CYP3A inhibitor according to claim 1, wherein said CYP3A inhibitor is orally administered to patients.

Claim 6. (Currently Amended) A pharmaceutical composition comprising the ~~The~~ CYP3A inhibitor according to claim 5 and, further comprising at least one pharmaceutically acceptable excipient ~~excipients~~.

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Claim 7. (Previously Presented) The CYP3A inhibitor according to claim 1, wherein said CYP3A inhibitor is administered to patients via food or in the form of capsule or tablet.

Claim 8. (Previously Presented) The CYP3A inhibitor according to claim 1, wherein said CYP3A inhibitor is co-administered with a first-pass effect drug.

Claim 9. (Previously Presented) The CYP3A inhibitor according to claim 9, wherein said first-pass effect drug and said CYP3A inhibitor are co-administered orally.

Claim 10. (Previously Presented) The CYP3A inhibitor according to claim 8, wherein said drug is one selected from the group consisting of erythromycin, felodipine, troleandomycin, nifedipine, cyclosporin, FK506, teffendine, tamoxifen, lidocaine, triazolam, dapsone, diltiazem, lovastatin, simvastatin, quinidine, ethylestradiol, testosterone, midazolam, and alfentanil.

Claim 11. (Previously Presented) The CYP3A inhibitor according to claim 8, wherein said CYP3A inhibitor is catechin, and wherein said first-pass effect drug is simvastatin.

Claim 12. (Previously Presented) The CYP3A inhibitor according to claim 1, wherein said CYP3A inhibitor is orally administered to patients with cancer.

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Claim 13. (Previously Presented) The CYP3A inhibitor according to claim 12, wherein said CYP3A cancer is intestinal or hepatic cancer.

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Claim 14. (Previously Presented) The CYP3A inhibitor according to claim 13, wherein said intestinal cancer is adenocarcinoma.

Claim 15. (Previously Presented) The CYP3A inhibitor according to claim 13, wherein said hepatic cancer is hepatoma.

Claim 16. (Withdrawn) A method for treating patient with intestinal or hepatic cancer comprising orally administering the CYP3A inhibitor according to claim 1 to said patient with intestinal or hepatic cancer.

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Claim 17. (Withdrawn) A cytochrome P450 3A (CYP3A) enhancer which is a free base or pharmacologically acceptable salt of at least one compound selected from the group consisting of apigenin, formononetin, and luteolin-7-glycoside.

Claim 18. (Withdrawn) The CYP3A enhancer according to claim 16, wherein said CYP3A enhancer induce the CYP3A enzymatic activity.

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Claim 19. (Withdrawn) A method for treating patients with hepatic failure comprising: treating said patients with hepatic failure with a CYP3A enhancer.

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Claim 20. (New) A method for prolonging a therapeutic effect of an orally administered drug in a mammal comprising orally administering a cytochrome P450 3A (CYP3A) inhibitor to said mammal;

wherein said orally administered drug is at least one selected from the group consisting of erythromycin, troleandomycin, teffendine, tamoxifen, lidocaine, triazolam, dapsone, diltiazem, lovastatin, simvastatin, quinidine, midazolam, and alfentanil; and

wherein said CYP3A inhibitor is at least one selected from the group consisting of α -naphthoflavone, β -naphthoflavone, apigenin, baicalein, β -myrcene, catechin, 3-phenylpropyl acetate, formononetin, hesperetin, hesperidin, isoquercitrin, lauryl alcohol, luteolin, luteolin-7-glycoside, narigin, nordihydroguaiaretic acid, quercitrin, swertiamarin, terpineol, and trans-cinnamaldehyde.

? support in spec.

Claim 21. (New) The method according to claim 20, wherein said CYP3A inhibitor is at least one selected from the group consisting of α -naphthoflavone, β -naphthoflavone, baicalein, catechin, 3-phenylpropyl acetate, formononetin, lauryl alcohol, luteolin, luteolin-7-glycoside, nordihydroguaiaretic acid, and swertiamarin.

Claim 22. (New) The method according to claim 20, wherein said orally administered drug and said CYP3A inhibitor are orally co-administered to said mammal.

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Claim 23. (New) The method according to claim 20, wherein said CYP3A inhibitor is catechin, and wherein said orally administered drug is simvastatin.

Claim 24. (New) A method for treating a patient suffered from intestinal or hepatic cancer comprising orally administering said patient with a cytochrome P450 3A (CYP3A) inhibitor, wherein said CYP3A inhibitor is at least one selected from the group consisting of α -naphthoflavone, β -naphthoflavone, apigenin, baicalein, β -myrcene, catechin, 3-phenylpropyl acetate, formononetin, gallic acid, hesperetin, hesperidin, isoquercitrin, lauryl alcohol, luteolin, luteolin-7-glycoside, narigin, nordihydroguaiaretic acid, quercitrin, swertiamarin, terpineol, and trans-cinnamaldehyde.

12
m >
Claim 25. (New) The method according to claim 25, wherein said CYP3A inhibitor is at least one selected from the group consisting of α -naphthoflavone, β -naphthoflavone, baicalein, catechin, 3-phenylpropyl acetate, formononetin, lauryl alcohol, luteolin, luteolin-7-glycoside, nordihydroguaiaretic acid, and swertiamarin.
